

THE UNITED STATES DISTRICT COURT FOR THE  
NORTHERN DISTRICT OF OKLAHOMA

W. A. DREW EDMONDSON, in his )  
capacity as ATTORNEY GENERAL )  
OF THE STATE OF OKLAHOMA and )  
OKLAHOMA SECRETARY OF THE )  
ENVIRONMENT C. MILES TOLBERT, )  
in his capacity as the )  
TRUSTEE FOR NATURAL RESOURCES )  
FOR THE STATE OF OKLAHOMA, )

Plaintiff, )

vs. )

TYSON FOODS, INC., et al, )

Defendants. )

4:05-CV-00329-TCK-SAJ

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MOTION FOR  
PRELIMINARY INJUNCTION HEARING  
  
BEFORE THE HONORABLE GREGORY FRIZZELL

VOLUME VII  
Daily Copy Transcript

March 10, 2008

1 A I do recall reading that, yes.

2 Q Now, as we're talking about microbial source  
3 tracking, is there a distinction to be made between  
4 the tools and the methodology?

5 A Well, you can make that distinction, yes.

02:59PM

6 Q Please tell us what, if any, distinction you  
7 would make.

8 A Well, the tools, I mean, I guess I think of  
9 the tools as your common laboratory procedures, your  
10 PCR, your cloning, your DNA sequencing, those kind  
11 of things we do in the lab every day. Those will  
12 put you to sleep. It's pretty boring stuff. It's  
13 how you use these tools that really --

02:59PM

14 Q It's the application of the tools?

15 A Exactly. It's the application, and in these,  
16 especially the molecular cases, what you're looking  
17 at, what piece of fragment, what bacteria. You  
18 know, this is really not even an identified  
19 bacteria. It's 98 percent genetically close to  
20 Brevibacteria avium. It's never been cultured.

03:00PM

21 When you find a bacteria, that's the first step you  
22 do.

03:00PM

23 Q Did Dr. Harwood culture it?

24 A I don't believe she did. Brevibacterium,  
25 actually to differentiate, there's Brevibacteria

03:00PM

1 Q Usually particle size, the smaller the  
2 particle size, the easier for the transport in the  
3 environment; isn't that a general truth, sir?

4 A It depends on the conditions of the field.  
5 It's a vegetation.

04:31PM

6 Q Okay. Now, isn't it true that Dr. Harwood did  
7 test all of those different locations except for  
8 poultry feces, poultry litter, land application,  
9 field surface, runoff waters, soils, the surface  
10 transport water, the groundwater and the ultimate  
11 recreational waters, and they found the poultry  
12 specific DNA in all those locations?

04:31PM

13 A I believe that they did find the  
14 Brevibacteria, that small strand, in all those  
15 locations. They did test -- you're correct, they  
16 tested all those.

04:31PM

17 Q I want to turn to Exhibit 40.

18 A Okay.

19 Q That was where you had the volume of --

20 MR. PAGE: Again, Your Honor, I know I step  
21 on very close -- but this is part of the cross, and  
22 we're going to talk a little bit more about cows but  
23 just briefly.

04:32PM

24 THE COURT: I understand.

25 MR. PAGE: Thank you.

04:32PM

1 primers that she produced caused replication. In  
2 many cases -- that's how you have to optimize and  
3 validate them. In many cases you have 19 bases that  
4 match on a 20 base primer, and you can get  
5 replication. I'm sorry not I'm answering yes or no  
6 but it's simply not that straightforward.

04:39PM

7 Q Would you look at State's 569? I'll represent  
8 to you, sir, that it's just a portion, actually  
9 attachments to your report.

10 A Yes, sir.

04:40PM

11 Q This is your work product; correct?

12 A I believe it is.

13 Q And under litter samples it says,  
14 Brevibacterium nanograms per gram on the first entry  
15 there; correct?

04:40PM

16 A Yes, it does.

17 Q What is that; what do you intend that column  
18 to represent?

19 A Well, that was the values that were reported  
20 in terms of the amount of Brevibacterium DNA that  
21 they, Dr. Harwood and North Wind, included in their  
22 data.

04:40PM

23 Q If that is not the amount of Brevibacterium  
24 that was reported, then your analysis and  
25 correlation would be mistaken; is that correct?

04:40PM

Page 2090

1 A It may be off a little. I don't know that it  
2 would affect the conclusions.

3 Q Let's take a look at it. Would you please  
4 look at the North Wind report dated October 4,  
5 State's Exhibit 533?

04:41PM

6 A Okay.

7 Q And -- well, first is of all, look at Page 4.

8 A Yep, I'm there.

9 Q I just want to correct the record. When you  
10 were looking at the flow chart for Dr. Harwood, you  
11 testified that she was mistaken on this chart, that  
12 the detection limit was more like 2,000 rather than  
13 6 gene copies; correct?

04:41PM

14 A I repeated what she said under oath, I thought  
15 that it was 2,000 either in her deposition or here.

04:41PM

16 Q Have you seen this North Wind report that's  
17 part of the evaluation for the PCR, the October 4th  
18 report?

19 A It was amended in December, sir.

20 Q There was additional reports issued in  
21 December, but this is one of the earliest reports  
22 that hasn't been amended. Did you not understand  
23 that --

04:41PM

24 MR. JORGENSEN: I object. Counsel's  
25 statement about which reports there are or what

04:42PM

1 dates they came, there's been no foundation for  
2 that.

3 THE COURT: Were you contesting, Mr.  
4 Jorgensen, that this report was not amended?

5 MR. JORGENSEN: I honestly don't know what  
6 days they came and which ones were amended. I don't  
7 think he does either, and counsel can't provide that  
8 testimony. Sorry, Your Honor.

04:42PM

9 THE COURT: In efforts to speed this up,  
10 Mr. Page is making representation to this witness.  
11 If you find to the contrary, you can bring it up.  
12 The objection is overruled. Mr. Page?

04:42PM

13 Q You see there on Page 4 where it says  
14 detection of poultry specific brevi biomarker, it  
15 says the detection limit is actually 6 copies per  
16 microliter?

04:42PM

17 A And I believe Dr. Harwood indicated that that  
18 detection was actually for the regular PCR, but for  
19 the quantitative PCR, because of the dilution steps,  
20 that she testified that it was 2,000 for the  
21 quantitative PCR.

04:43PM

22 Q Okay. So it's 2,000 quantitative and 6 for  
23 detect or non-detect?

24 A Correct.

25 Q Present or not?

04:43PM

Page 2092

1 A Exactly.

2 Q Now, I want you to look on that same exhibit  
3 to the page -- two more pages beyond where it has a  
4 list of the results.

5 A Yep.

04:43PM

6 Q Now, under Exhibit 569, you've listed values  
7 of 21, 21.3. Do you see those numbers there from  
8 569?

9 A I do.

10 Q Now, under what column on Exhibit 533 are  
11 those levels of DNA located?

04:43PM

12 A Those are under the DNA.

13 Q So that's total DNA, is it not, sir?

14 A That would be.

15 Q So you made a mistake when you did your  
16 correlation analysis?

04:44PM

17 A I'd have to double check. I don't know if  
18 this -- what was shown earlier was based on the same  
19 data.

20 Q Well, I mean, we could look at it. Look at  
21 your Defendant's Exhibit D 42.

04:44PM

22 A It might have been, sir.

23 Q Don't you see the same plots there for total  
24 DNA rather than individual strands of Brevibacteria?

25 A Yes, sir, I do.

04:44PM

1 Q Okay. So if you use the proper correlation  
2 analysis, is it possible that this might actually  
3 show a correlation between the litter and an  
4 indicator bacteria?

5 A I actually did use these numbers on the right  
6 as well. Where I got the DNA in the first column --  
7 or I'm sorry -- the second column here where it says  
8 nanograms per liter, the DNA was the database that  
9 was provided to me by the State on an Excel  
10 spreadsheet that had these numbers here listed as  
11 the qPCR.

04:44PM

04:45PM

12 Q You also had the October 4th report, did you  
13 not?

14 A The October 4th, I probably did.

15 Q And that is very clear that the numbers you  
16 used were total DNA rather than biomarker copies per  
17 microliter; correct?

04:45PM

18 A In this but again --

19 Q Can you answer that yes or no, please?

20 A I didn't base -- I didn't get that from this  
21 report. So when I did my analysis, I'm sorry, I  
22 didn't base it on this document.

04:45PM

23 Q You didn't go to the original source?

24 A Well, I was looking at more recent documents.  
25 We had the North Wind report in November, December.

04:45PM



1 Some of the DNA were negative values on some of the  
2 reports I'd seen. So it was very dynamic, not only  
3 in the column headings, but in the numbers.

4 Q In your Exhibit 42, when you say biomarker,  
5 nanograms per gram under the horizontal line, the  
6 base line there, that's really a mistake; that's not  
7 the biomarker; that's total DNA; correct?

04:46PM

8 A It potentially may be, sir.

9 Q Would you please turn to State's Exhibit 534,  
10 please?

04:46PM

11 A Okay.

12 Q I'll represent to you that this is a  
13 correlation plot using the actual gene copies from  
14 the October 7th and comparing it to Enterococcus.

15 Do you see a correlation on State's Exhibit 534?

04:46PM

16 A There would be -- it looks like with respect  
17 to Enterococcus it is a correlation.

18 Q Thank you, sir. Now, isn't it true that for  
19 Enterococcus, that particular indicator bacteria has  
20 been referenced by the State as causing a  
21 significant amount of exceedances in the state water  
22 quality?

04:47PM

23 A I think about 5,800 miles in the IRW -- or I'm  
24 sorry -- in the state are impaired, listed as  
25 impaired by Enterococci.

04:47PM